

10-17-02
1653
EXPRESS MAIL NO.: EL 451 598 589 US

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

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OCT 22 2002

TECH CENTER 1600/2900

Application of : Dasseux, et al.

Serial No. : 09/865,989

Group Art Unit: 1653

Filed: May 25, 2001

Examiner: Russell

For: APOLIPOPROTEIN A-I
AGONISTS AND THEIR
USE TO TREAT
DYSLIPIDEMIC DISORDERS

Attorney Docket No.: 9196-019-999

FEE TRANSMITTAL SHEETCommissioner for Patents
Washington, D.C. 20231

Sir:

The fee required to be filed with the accompanying amendment of even date herewith concerning the above-identified application has been estimated to be \$54.

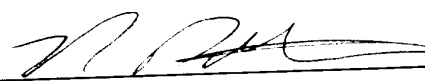
The claim amendment fee has been estimated as shown below:

(Col. 1)			(Col. 2)			(Col. 3)			SMALL ENTITY			OTHER THAN A SMALL ENTITY						
CLAIMS REMAINING AFTER AMENDMENT			HIGHEST NO PREVIOUSLY PAID FOR			PRESENT EXTRA			RATE		ADDIT. FEE		OR		RATE		ADDIT. FEE	
TOTAL	46	MINUS	40	=	6	×	9	\$	54.00	×	18	\$						
INDEP.	01	MINUS	05	=	0	×	42	\$	0.00	×	84	\$						
<input type="checkbox"/> FIRST PRESENTATION OF MULTIPLE DEP. CLAIM									140	\$			280	\$				
									TOTAL	\$	54.00	OR	TOTAL	\$				

Please charge the required fee to Pennie & Edmonds LLP Deposit Account No. 16-1150. A copy of this sheet is enclosed.

Respectfully submitted,

Date: October 16, 2002


42,983
(Reg No.)
Rahul Pathak
for Laura A. Coruzzi (Reg. No. 30,742)
PENNIE & EDMONDS LLP
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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

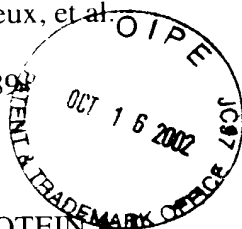
RECEIVED

Application of : Dasseux, et al.

Serial No. : 09/865,989

Filed: May 25, 2001

For: APOLIPOPROTEIN A-1
AGONISTS AND THEIR
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DYSLIPIDEMIC DISORDERS



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Examiner: Russell

Attorney Docket No.:
9196-019-999

OCT 22 2002

TECH CENTER 1600/2900

PRELIMINARY AMENDMENT UNDER 37 C.F.R. §1.115

Commissioner for Patents
Washington, D.C. 20231

Sir:

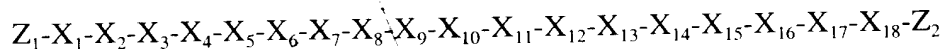
In accordance with 37 C.F.R. §115, please enter the following amendments prior to examination of the above-captioned application.

AMENDMENTS

IN THE CLAIMS

Please amend Claims 1, 16, 17, 18, 25 and 33 as follows:

1. (Twice amended) An ApoA-I agonist compound comprising:
(i) an 18 to 22-residue D-enantiomeric peptide or peptide analogue which forms an amphipathic α -helix in the presence of lipids and which comprises formula (I):



X_1 is D-Ala (a), Gly (G), D-Asn (n), D-Gln (q) or D-Pro (p);

X_2 is a D-enantiomeric aliphatic residue;

X_3 is D-Leu (l);

X_4 is a D-enantiomeric acidic residue;

X_5 is D-Leu (l) or D-Phe (f);